

FORM PTO-1390 (Modified)
(REV 11-98)

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

TRANSMITTAL LETTER TO THE UNITED STATES

DESIGNATED/ELECTED OFFICE (DO/EO/US)

CONCERNING A FILING UNDER 35 U.S.C. 371

228.1007

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR

60/086,895 09/701450

INTERNATIONAL APPLICATION NO.

PCT/EP99/03681

INTERNATIONAL FILING DATE

27 May 1999

PRIORITY DATE CLAIMED

27 May 1998

TITLE OF INVENTION

PREPARATIONS FOR THE APPLICATION OF ANTI-INFLAMMATORY, ESPECIALLY ANTISEPTIC AGENTS AND/OR AGENTS PROMOTING THE HEALING OF WOUNDS, TO THE LOWER RESPIRATORY TRACT

APPLICANT(S) FOR DO/EO/US

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371 (c) (2))
 - a. ☒ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ A copy of the International Search Report (PCT/ISA/210).
8. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3))
 - a. ☒ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☐ have not been made and will not be made.
9. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
10. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)).
11. ☒ A copy of the International Preliminary Examination Report (PCT/IPEA/409).
12. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)).

Items 13 to 20 below concern document(s) or information included:

13. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
14. ☒ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
15. ☒ A **FIRST** preliminary amendment.
16. ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
17. ☐ A substitute specification.
18. ☐ A change of power of attorney and/or address letter.
19. ☒ Certificate of Mailing by Express Mail
20. ☒ Other items or information:

- Letter re: Priority
- Response to Written Opinion
- Postcard
- Abstract on separate sheet
- Written Opinion

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR 1.53) 60/086,895	INTERNATIONAL APPLICATION NO. 09/701450	ATTORNEY'S DOCKET NUMBER 228.1007
PCT/EP99/03681		

21. The following fees are submitted:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :

- ☐ Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2) paid to USPTO and International Search Report not prepared by the EPO or JPO **\$1,000.00**
- ☒ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO **\$860.00**
- ☐ International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO **\$710.00**
- ☐ International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4) **\$690.00**
- ☐ International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4) **\$100.00**

ENTER APPROPRIATE BASIC FEE AMOUNT =**\$860.00**

Surcharge of **\$130.00** for furnishing the oath or declaration later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (e)).

\$0.00

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE		
Total claims	75 - 20 =	55	x \$18.00	\$990.00	
Independent claims	3 - 3 =	0	x \$80.00	\$0.00	

Multiple Dependent Claims (check if applicable).

☒**\$270.00****TOTAL OF ABOVE CALCULATIONS =****\$2,120.00**

Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28) (check if applicable).

☐**\$0.00****SUBTOTAL =****\$2,120.00**

Processing fee of **\$130.00** for furnishing the English translation later than ☐ 20 ☐ 30 months from the earliest claimed priority date (37 CFR 1.492 (f)).

+

\$0.00**TOTAL NATIONAL FEE =****\$2,120.00**

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable).

☐**\$0.00****TOTAL FEES ENCLOSED =****\$2,120.00**

Amount to be: refunded	\$
charged	\$

☒ A check in the amount of **\$2,120.00** to cover the above fees is enclosed.

☐ Please charge my Deposit Account No. _____ in the amount of _____ to cover the above fees.
A duplicate copy of this sheet is enclosed.

☒ The Commissioner is hereby authorized to charge any fees which may be required, or credit any overpayment to Deposit Account No. **50-0552** A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

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Morey B. Wildes

NAME

39,968

REGISTRATION NUMBER

November 27, 2000

DATE

228.1007

UNITED STATES PATENT & TRADEMARK OFFICE

Re: Application of: Wolfgang FLEISCHER and Karin REIMER
Serial No.: To Be Assigned
Filed: Simultaneously Herewith
For: **PREPARATIONS FOR THE APPLICATION OF
ANTI-INFLAMMATORY, ESPECIALLY
ANTISEPTIC AGENTS AND/OR AGENTS
PROMOTING THE HEALING OF WOUNDS, TO
THE LOWER RESPIRATORY TRACT**

PRELIMINARY AMENDMENT

Box: PCT
Assistant Commissioner for Patents
Washington, D.C. 20231

November 27, 2000

Sir:

Prior to examining the application, kindly amend the original published claims as follows:

IN THE ABSTRACT

Please **add** the enclosed abstract numbered as a separate sheet.

IN THE CLAIMS

Please **amend** the claims as follows:

1. (Amended) A **[process for the manufacture of a]** pharmaceutical preparation for the application of antiseptic agents **[and/or] or** agents which promote the healing of wounds to the lower respiratory tract, **[characterized in that] wherein** the preparation contains at least one of

said agents combined with a particulate carrier.

2. (Amended) The **preparation [process]** of claim 1, **wherein** [characterised in that] said particulate carrier comprises at least one of a liposome preparation, a microsphere preparation, a nanoparticle preparation, a Large Porous Particle preparation, or a laser-pulse polymer coated molecule preparation.

3. (Amended) The **preparation [process according to claim 1 or 2] of claim 1, wherein** [characterised in that] at least the greatest part of said agent is encapsulated inside the carrier, especially a liposome or microsphere carrier.

4. (Amended) The **preparation [process of any one of claims 1 to 3] of claim 1, wherein** [characterised in that] the antiseptic agent is selected from oxygen-**releasing compounds**, and halogen-releasing compounds, metal compounds, such as silver and mercury compounds; organic disinfectants including inter alia formaldehyde-releasing compounds, alcohols, phenols, including alkyl**phenols**, [- and] arylphenols as well as halogenated phenols, quinolines and acridines, hexahydropyrimidines, quaternary ammonium compounds and iminium salts, and guanidines.

5. (Amended) The **preparation [process according to] of claim 4, wherein** [characterised in that] the antiseptic agent is selected from the group comprising metal compounds such as

mercury compounds, phenol derivatives such as thymol, eugenol and hexachlorophene, iodine and iodine complexes.

6. (Amended) The **preparation of** [process according to] claim 5, **wherein** [characterised in that] the antiseptic agent is povidone iodine.

7. (Amended) The **preparation** [process according to any one of claims 1 to 6] **of claim 1,** **wherein** [characterised in that] the wound-healing promoting agent is selected from agents promoting granulation and epithelization such as dexpanthenol, allantoines, azulenes, tannines, compounds from the vitamin B series, or similarly acting agents.

8. (Amended) The **preparation** [process according to any one of the preceding claims] **of claim 1, wherein** [characterised in that] the preparation contains at least one antiseptic and at least one wound-healing promoting agent.

9. (Amended) The **preparation** [process according to any one of the preceding claims] **of claim 1, wherein** [characterized in that] the carrier particles, [especially liposomes,] have a substantially uniform size in the range between about 1 μm and about 50 μm [, preferably in the range between about 1 and about 30 μm].

10. (Amended) The **preparation of claim 9** [process according to claim 9] **wherein,**

[characterised in that] the carrier particles, [especially liposomes,] have a substantially uniform size in the range between 20 μm and 30 μm diameter for application to the trachea[, in the range between about 10 and 20 μm diameter for application to the bronchi and between about 1 and 6 μm , especially between 2 and 5 μm , diameter for application to the alveoli].

11. (Amended) The preparation [process according to any of the preceding claims] of claim 1, wherein [characterised in that] the carrier[, especially liposome, preparation] releases the agent over an extended time period[, preferably an extended time period of several hours duration].

12. (Amended) The preparation of [process according to] claim 11, wherein [characterised in that] the carrier[, especially liposome, preparation] releases the agent at approximately the same release rate over the release time period.

13. (Amended) The preparation of [process according to any one of the preceding claims] claim 1, wherein [characterised in that] the preparation additionally comprises at least one anesthetically active agent.

14. (Amended) The preparation [process according to any one of the preceding claims] of claim 1, wherein [characterized in that] the preparation contains additives and adjuvants [such as] comprising conserving agents, antioxidants and consistency-forming additives.

15. (Amended) The preparation [process according to any one of claims 1 to 14] of claim 1, wherein the preparation [being] comprises a suitable form for administration via the lower respiratory tract comprising [the] an active-agent loaded carrier, wherein the carrier is [especially] in the form of liposomes, [preferably] in the form of an aerosol, or [especially] in the form of a powder aerosol.

16. (Amended) The preparation [process according to any one of claims 1 to 14] of claim 1, wherein the preparation [being in the form of] comprises a compacted solid medicament reservoir, [preferably] a ring-tablet, [more preferably] a gelatin capsule, a powder, a spray, an emulsion, a dispersion, a suspension or a solution containing the carrier and agent or agents in a pharmaceutically acceptable solid or liquid formulation, which is suitable for the generation of inhalable particles.

17. (Amended) The preparation [process according to any one of the preceding claims] of claim 1, [being in] comprising a suitable form for administration via the lower respiratory tract, which comprises:

(a) liposomes comprising a pharmaceutically acceptable liposome membrane forming substance; and

(b) a 0.1 to 2% PVP iodine solution (at approximately 10% available iodine in the PVP iodine complex) at least most of which is encapsulated by said liposome membranes,

wherein the liposomes are of substantially uniform size between about 1 μm and about 50

μm , and[, **in case,**] the formulation additionally comprises customary additives, adjuvants and auxiliary substances of a pharmaceutical solution or dispersion formulation.

18. (Amended) The **preparation [process]** according to claim 17, characterised in that the liposomes are of a substantially uniform size, in the range between 20 **μm** and 30 μm diameter for application to the trachea[, **in the range between about 10 and 20 μm diameter for application to the bronchi and between about 1 and 6 μm , especially between 2 and 5 μm , diameter for application to the alveoli].**

19. (Amended) The **preparation [process according to any one of claims 1 to 18] of claim 1**, wherein the preparation is suited for the treatment of infectious diseases or alleviation of diseases such as HIV infections which are accompanied by opportunistic infections or a suppressed immune system.

20. (Amended) The **preparation [process according to any one of claims 1 to 18] of claim 1**, wherein the preparation is suited for the treatment of acute **bronchitis**, [and/or] chronic bronchitis, pneumonia, bronchiectasia, cystic fibrosis, diphtheria [and/or] **or** tuberculosis.

21. (Amended) The **preparation [process according to any one of claims 1 to 20] of claim 1**, wherein the preparation is suited for functional and cosmetic tissue remodeling and repair treatments.

22. (Amended) A method of preventing or treating infections of the human or animal lower respiratory tract **comprising** [, by] applying, to said tract, a pharmaceutical preparation comprising at least one antiseptic agent [**and/or**] **or** wound-healing promoting agent, [**said agent being**] combined with a particulate carrier in said preparation.

23. (Amended) A method of functional and cosmetic tissue remodeling and repair in the human or animal lower respiratory tract [, by] **comprising** applying, to said tract, a pharmaceutical preparation comprising at least one anti-inflammatory, [**especially**] antiseptic [**and/or**] **or** wound-healing promoting agent combined with a [**particulatar**] **particulate** carrier.

26. (Amended) The method of claim 23, wherein the anti-inflammatory agent is selected from antiseptic agents, antibiotics, corticosteroids [**and**] **or** wound-healing promoting agents.

27. (Amended) The method of claim 22 or 23, wherein the antiseptic agent is selected from oxygen-**releasing compounds** and halogen-releasing compounds; metal compounds, such as silver **compounds** and mercury compounds; organic disinfectants including inter alia formaldehyde-releasing compounds, alcohols, phenols including alkyl**phenyols**[- **and**] arylphenols as well as halogenated phenols, quinolines and acridines, hexahydropyrimidines, quaternary ammonium compounds and iminium salts, and guanidines.

32. (Amended) The method of claim 22 or 23, wherein the carrier particles[, especially

liposomes,] have a substantially uniform size in the range between about 1 μm and about 50 μm], **preferably in the range about 1 μm and about 30 μm**].

33. (Amended) The method according to claim 32, wherein the carrier particles[, **especially liposomes,**] have a substantially uniform size in the range between 20 μm and 30 μm diameter for application to the trachea[, **in the range between about 10 and 20 μm diameter for application to the bronchi and between about 1 and 6 μm diameter, especially between 2 and 5 μm , for application to the alveoli**].

34. (Amended) The method of claim 22 or 23, wherein that the carrier[, **especially liposome, preparation**] releases the agent over an extended time period[, **preferably an extended time period of several hours duration**].

35. (Amended) The method of claim 22 or 23, wherein the carrier[, **especially liposome, preparation**] releases the agent at approximately the same release rate over the release time period.

38. (Amended) The method of claim 22 or 23, **wherein** the preparation [being in] **comprises** a suitable form for administration via the lower respiratory tract, comprising the active-agent loaded carrier, [**especially**] in the form of liposomes, [**preferably**] in the form of an aerosol, [**especially**] **or** in the form of a powder aerosol.

39. (Amended) The method of claim 22 or 23, **wherein** the preparation **[being in the form of] comprises** a compacted solid medicament reservoir, **[preferably]** a ring-tablet, **[more preferably]** a gelatine capsule, a powder, a spray, an emulsion, a dispersion, a suspension or a solution containing the carrier and agent or agents in a pharmaceutically acceptable solid or liquid formulation, which is suitable for the generation of inhalable particles.

41. (Amended) The method of claim 22 or 23, wherein the liposomes are of a substantially uniform size in the range between 20 μm and 30 μm diameter for application to the trachea **[, in the range between about 10 and 20 μm diameter for application to the bronchi and between about 1 and 6 μm , especially between 2 and 5 μm , diameter for application to the alveoli].**

43. (Amended) The method of claim 22 or 23, wherein the preparation is suited for the treatment of acute **bronchitis**, **[and/or] or** chronic bronchitis, pneumonia, bronchiectasia, cystic fibrosis, diphtheria **[and/or] or** tuberculosis.

Please **add** the new claims as follows:

--44. (New) The preparation according to claim 9, wherein the carrier particles, have a substantially uniform size in the range between about 1 μm about 30 μm .---

--45. (New) The preparation according to claim 10, wherein, the carrier particles, have a

substantially uniform size in the range between about 10 μm and 20 μm diameter for application to the bronchi.--

--46. (New) The preparation according to claim 10, wherein, the carrier particles, have a substantially uniform size in the range between about 1 μm and 6 μm for application to the alveoli.--

--47. (New) The preparation according to claim 10, wherein, the carrier particles, have a substantially uniform size in the range between about 2 μm and 5 μm for application to the alveoli.--

--48. (New) The preparation according to claim 17, wherein the liposomes are of a substantially uniform size, in the range between 10 μm and 20 μm diameter for application to the bronchi.--

--49. (New) The preparation according to claim 17, wherein the liposomes are of a substantially uniform size, in the range between 1 μm and 6 μm diameter for application to the alveoli.--

--50. (New) The preparation according to claim 17, wherein the liposomes are of a substantially uniform size, in the range between 2 μm and 5 μm diameter for application to the alveoli.-

- 51. (New) The method of claim 22 or 23, wherein the carrier particles have a substantially uniform size in the range between about 1 μm and about 30 μm .--
- 52. (New) The method according to claim 32, wherein the carrier particles have a substantially uniform size in the range between 10 μm and 20 μm diameter for application to the bronchi.--
- 53. (New) The method according to claim 32, wherein the carrier particles have a substantially uniform size in the range between 1 μm and 6 μm diameter for application to the alveoli.--
- 54. (New) The method according to claim 32, wherein the carrier particles have a substantially uniform size in the range between 2 μm and 5 μm diameter for application to the alveoli.--
- 55. (New) The method of claim 22 or 23, wherein the liposomes are of a substantially uniform size in the range between 10 μm and 20 μm diameter for application to the bronchi.--
- 56. (New) The method of claim 55, wherein the liposomes are of a substantially uniform size in the range between 1 μm and 6 μm diameter for application to the alveoli.--
- 57. (New) The method of claim 56, wherein the liposomes are of a substantially uniform size

in the range between 2 μm and 5 μm diameter for application to the alveoli.--

REMARKS

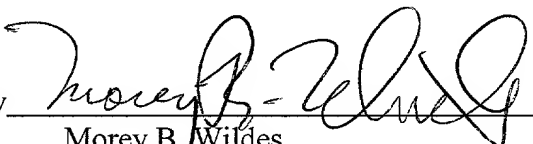
Entry of the above amendments to the claims is respectfully requested. Applicants have amended the claims in order to conform the claims U.S. patent practice. No new matter has been added by way of these amendments.

Further, the applicants have enclosed herewith an abstract on a separate sheet.

The Assistant commissioner is authorized to charge any additional fees to Attorney Deposit Account No. 50-0552.

Respectfully submitted,

DAVIDSON, DAVIDSON & KAPPEL, LLC

By 
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ABSTRACT

Use of an anti-inflammatory agent such as povidine iodine for the preparation of a pharmaceutical composition for the treatment of diseases of the lower respiratory tract which are susceptible to the administration of such agents.

Preparations for the application of anti-inflammatory, especially
antiseptic agents and/or agents promoting the healing of wounds,
to the lower respiratory tract

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The invention concerns preparations for the application of agents with anti-inflammatory, especially antiseptic and/or wound healing promoting properties to the lower respiratory tract. The preparations are specifically applied to trachea, bronchi and alveoli in the lower respiratory tracts of humans and animals.

10

Furthermore, the invention concerns a method of preventing or treating infections by applying a pharmaceutical preparation.

15

A plurality of different antibiotic and antiseptic agents are known for the topical treatment of infectious maladies. A decisive disadvantage of antibiotic agents is that the infecting bacteria show primary resistances, and can acquire secondary resistances, against these agents. Further, antibiotics quite often lead to patient sensibilisation. The use of e.g. halogen-releasing antiseptics such as povidone iodine, also known as polyvidone iodine or PVP-iodine, i.e. the poly(1-vinyl-2-pyrrolidin-2-one)-iodine complex, can prevent resistances. Antiseptic agents are also much more rarely allergenic as compared to antibiotics.

20

At present, infectious diseases of the respiratory tract are treated with antibiotics. The application of antibiotic agents via the respiratory tract has been the subject of several reviews and articles with an emphasis on the lower respiratory tract. Ramsey et al., for example, describe the intermittent administration of inhaled tobramycin in patients with cystic fibrosis in "The New England Journal of Medicine", Volume 340, Number 1, 1999, p. 23-30.

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The aerosolization of imipenem/cilastatin for preventing pseudomonas-induced acute lung injury has been investigated by Wiener-Kronish in "Journal of Antimicrobiol Chemotherapy" (1996) 38, p. 809-818.

5 Pulmonary applications of different antibiotic agents, like benzyl penicillin, tobramycin or amikacin, for the treatment of infectious diseases are described by Schreier in several recent reviews, e.g. in "Medical applications of liposomes", Papahadjopoulos and Lasic (eds.), Elsevier 1998.

10 However, the treatment with antibiotics leads to the complications known to the skilled person. For example, patients suffering from acute or chronic bronchitis are often treated with antibiotics in order to alleviate the symptoms. This often merely leads to resistances of the bacteria responsible for the symptoms. Many diseases of the respiratory tract are caused by viruses. Antibiotics are inefficient
15 in such cases, and such patients are not cured of the infections.

The use of antiseptics and/or wound-healing promoting agents for external application to humans and animals is disclosed in our earlier patent EP 0 639 373. Specifically, liposome preparations of PVP-iodine are shown therein to be
20 topically applicable to the external parts of the eye. These preparations generally take the form of a cream, an ointment, a lotion, a gel or a drop formulation.

Liposomes are well-known drug carriers and therefore the application of medicaments in liposomal form has been subject of investigation for quite some
25 time. An overview concerning pulmonary delivery of liposome encapsulated drugs in asthma therapy is provided by the review "Pulmonary delivery of liposomes" (H. Schreier, in "Journal of Controlled Release", 24, 1993, p.209-223). The physicochemical characterization of liposome aerosols and also their therapeutic applications to the respiratory tract are shown therein. Drugs that have

- 3 -

been investigated for pulmonary delivery via liposomes include, e.g. anti-cancer agents, peptides, enzymes, anti-asthmatic and anti-allergic compounds and, as mentioned above, also antibiotics. The formulation of liposome aerosols or liposome powder aerosols using, for example a dry powder inhaler has also been described by H.

Schreier in "Formulation and in vitro performance of liposome powder aerosols" (S.T.P. Pharma Sciences 4, 1994, p.38-44).

Although a lot of attention has been paid to liposomes as drug carriers, as can be seen from the cited documents, there appears to be no prior art relating to liposomes and other particulates as carriers of anti-inflammatory, antiseptic and/or wound-healing promoting agents for applications in the body, especially in the lower respiratory tract, including the trachea, bronchi and alveoli.

Some of the prior art cited above is concerned with liposome preparations. It should be understood that alternative drug carriers of a similarly particulate character exist. These drug carriers can often -and also in the context of this invention- be used instead of liposomes and include microspheres (generally comprising lipophilic polymers), nanoparticles, "Large Porous Particles" and individually coated drug substance molecules, e.g. made by using pulsed laser deposition (PLD) techniques. These PLD methods can be used to apply coatings to drug powders and to modify surface properties and release rate to a variety of drug systems.

Where hereinafter reference is made to liposomes or particulate carriers, it is to be understood that this is to incorporate such alternative carriers, too.

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It is known in the art that the administration of inhalable particles to the respiratory tract can be achieved by nebulization or aerosolization of the liposome, microsphere, Large Porous Particle, PLD or nanoparticle preparations or by dry powder inhalation of the respective preparation.

5

There appears to be a marked reluctance in the art, to apply disinfectants to interior parts of the body, except maybe in extreme cases of life-threatening septical complications.

10 Generally, antibiotic preparations appear to be preferred, even in view of their above-discussed disadvantages.

15 An object of the instant invention is to provide a well tolerated, easily applicable anti-inflammatory, antiseptic and/or wound-healing promoting preparation, which provides protracted release and protracted topical effect of the active agent in the lower respiratory tract.

20 According to the invention this object is attained in that the preparation comprises at least one anti-inflammatory, antiseptic and/or wound healing promoting agent in the form of a particulate carrier preparation, as defined in independent claim 1.

The invention further comprises a method of treating the lower respiratory tract, in humans and animals, as defined in independent claims 21 and 22.

25 The dependent claims define further advantageous embodiments of the invention.

In the context of this invention, anti-inflammatory agents are understood to include antiseptic agents, antibiotic agents, corticosteroids, and wound-healing agents, as defined below.

- 5 -

In the context of this invention, antiseptic agents are understood to include those disinfecting agents which are pharmaceutically acceptable and suitable for the treatment of the lower respiratory tract to the extent that they can be formulated in accordance with the invention.

5

More specifically, antiseptic agents include inter alia oxygen- and halogen-releasing compounds; metal compounds, e.g. silver and mercury compounds; organic disinfectants including inter alia formaldehyde-releasing compounds, alcohols, phenols including alkyl- and arylphenols as well as halogenated phenols, quinolines and acridines, hexahydropyrimidines, quaternary ammonium compounds and iminium salts, and guanidines.

10

Wound-healing agents comprise agents promoting granulation and epithelization such as dexpanthenol, allantoines, azulenes, tannines, and vitamine B-type compounds.

15

The invention is premised on the surprising fact that particulate carriers, especially liposomes, but also microspheres, nanoparticles and coated drug substance molecules, are highly suited as carriers for antiseptic agents, especially for povidone iodine, and for agents promoting the healing of wounds, for application to the lower respiratory tract.

20

The preparations according to this invention permit protracted release of the agent or agents, and provide an extended and topical activity at the desired locus of action by interaction with cell surfaces.

25

The invention is, another aspect, based on a further surprising and unexpected fact. It is well known in the art that the formation of new body tissues may cause problems. Thus, it is known that body tissue repair may be accompanied by the

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formation of scar tissue, which can be functionally and/or cosmetically harmful, or at least undesirable. Hyperkeratosis and the uncontrolled proliferation of tissue may cause serious harm, leading to dysfunctions, and may of course also be cosmetically undesirable. After infections and inflammations, re-growing or
5 healing tissue may cause neoplasms and intergrowth. It is thus well known in the art that in the curing of diseases, proper remodelling of tissue is not only desirable, but in fact necessary.

It has now been surprisingly found that the use of anti-inflammatory agents, singly
10 or in combination with other such agents, leads to markedly less formation of undesirable body tissue in the course of tissue repair and other tissue growth processes. Thus, the formation of scar tissues is reduced, in skin but also in mucosa and in other tissues, such as muscle or inner organ tissues. Hyperkeratosis may be entirely suppressed, and intergrowth, or neoplasm formation in the curing
15 of infective diseases is also highly reduced.

One object achieved by the invention is therefore concerned with improved tissue repair in the body. The invention achieves this by the application of anti-inflammatory agents, in the form of a particulate carrier preparation as defined in
20 the independent claims.

The anti-inflammatory, antiseptic and/or wound-healing preparation can be administered to the respiratory tract by a nebulization agent loaded of the particulate carrier preparation, or by dry powder inhalation of the respective
25 preparation. For example, a liposome preparation can be made by loading liposomes with PVP iodine in a conventional procedure.

It is also possible to compact the loaded liposomes, optionally together with auxiliary materials, such as low molecular sugars, preferably lactose, to a tightly

- 7 -

compacted solid medicament reservoir. This medicament stock can then be abraded or micronized or treated in other ways to yield the powder in particle form. The resulting liposome preparation can be administered by inhalation of the preparation in the form of a powder aerosol, as, for example, described in "Acute
5 Effects of Liposome Aerosol Inhalation on Pulmonary Function in Healthy Human Volunteers" (Thomas et al., Preliminary report, Volume 99, 1991, p. 1268-1270). The pressures for preparing the tightly compacted solid medicament stock are preferably in the range of from 50-500 MPa. Such medicament stock is described in WO 94/14490 and a device for administration is disclosed in WO 93/24165.

10 The nature or constitution of the liposomes is generally not critical. The liposome preparation as, for example, described in EP 0 639 373 can be administered by inhalation as an aerosol. The disclosure of EP 0 639 373 is incorporated by reference.

15 The preparations according to this invention apparently do not only contain the active agent, like povidone iodine, encapsulated in the particulate carrier, especially in liposomes. It seems that there is also some amount of agent which is not contained inside the carrier. The preparations according to the invention often
20 show a marked initial effect which is observed in addition to the slower, protracted release of the active agent from the carrier. This effect is especially observed where the carrier comprises liposomes. Without wishing to be bound to any theoretical explanation, it is presently assumed that in addition to active agent encapsulated inside the liposomes, some active agent is present outside of the
25 liposomes, and probably loosely bound to the outer surfaces of the liposomes. This could be due to association of active agent molecules with the liposomal membrane, or it could be due to active agent molecules forming a layer on the liposomal surface, which layer partly or even fully coats the liposome externally. The type and amount of this initial agent effect can e.g. be influenced by choice

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of the concentration parameters.

The amphiphilic substances generally known in prior art to form liposome membranes can be employed in the context of the invention as long as they are pharmaceutically acceptable for the intended application. Presently, liposome forming systems comprising lecithin are preferred. Such systems can comprise hydrogenated soy bean lecithin besides cholesterol and disodium succinate-hexahydrate; it is presently specifically preferred to use hydrogenated soy bean lecithin as the sole membrane-forming agent.

10

The known prior art methods for forming liposome structures are described in the documents cited above and can generally be used in the context of the invention.

Broadly, these methods comprise mechanical agitation of a suitable mixture containing the membrane forming substance and water or an aqueous solution.

15

Filtration through suitable membranes is preferred in forming a substantially uniform liposome size.

20

The average size of the liposomes according to this invention can vary over a broad range, generally from about 1 to about 50 μm , preferably in the range of 1 and 30 μm diameter. For solutions, smaller average diameters, e.g. diameters of about 100 nm, may be more suitable.

25

The liposomes according to this invention have a substantially uniform size in the range between about 20 and 30 μm diameter for application to the trachea, in the range between about 10 and 20 μm diameter for application to the bronchi and between about 1 and 6 μm , especially between 2 and 5 μm , diameter for application to the alveoli.

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Where alternative particulate carriers are used, they are generally prepared as known in the art. Thus, microspheres which are used to deliver a very wide range of therapeutic or cosmetic agents, are made as described for example in WO 95/15118.

5

Nanoparticles may in some cases be used, provided that they can be loaded with a sufficient amount of active agent and can be administered to the lower respiratory tract according to this invention. They can be prepared according to the methods known in the art, as e.g. described by Heyder (GSF München) in "Drugs delivered to the lung", Abstracts IV, Hilton Head Island Conference, May 1998.

10

Methods using a pulse laser deposition (PLD) apparatus and a polymeric target to apply coatings to drug powders in a short non-aqueous process are also suitable for the formation of particulate preparations according to this invention. These have e.g. been described by Talton et al., "Novel Coating Method for Improved Dry Delivery", Univ. of Florida UF 1887 (1998).

15

A further suitable delivery system employs Large Porous Particles as disclosed by David A. Edwards et al. in "Large Porous Particles for Pulmonary Drug Delivery" (Science, 20. June 1997, Vol. 276, p. 1868-1871). The average size of Large Porous Particles according to this invention can e.g. be in the range of between about 5 and 20 μm diameter for application to the alveoli.

20

Preferred anti-inflammatory agents comprise antiseptic agents, antibiotics, corticosteroids and wound-healing promoting agents, as single substances or in combination with each other.

25

Preferred antiseptic agents comprise the well-known pharmaceutical substances providing fast effect, a broad range of activity, low systemic toxicity and good

- 10 -

tissue compatibility. They can e.g. be selected from the group comprising metal compounds, phenolic compounds, detergents, iodine and iodine complexes. A specifically preferred antiseptic agent is povidone iodine.

- 5 Preferred agents promoting the healing of wounds comprise substances which have been described in the literature for such application. Preferred such agents include substances known to promote epithelisation. These include vitamins, specifically from the vitamin B group, allantoin, some azulenes etc.
- 10 Some presently highly preferred embodiments of the invention comprise anti-inflammatory agents or combinations of such agents which show beneficial effects in tissue repair, especially with respect to functional and cosmetic tissue remodelling. In these embodiments, the active agent is often an antiseptic, such as PVP-iodine, or an antibiotic.
- 15 In preferred embodiments, the invention's preparations containing anti-inflammatory, especially antiseptic and/or wound-healing promoting agents can comprise further agents such as anaesthetic agents. Inventive preparations can also contain customary further agents, including adjuvants and additives,
- 20 antioxidants, conserving agents or consistency-forming agents such as viscosity adjusting additives, emulgators etc.
- 25 Generally, the concentrations in the preparation, particle sizes, active agent loadings etc. will be selected for such alternative carriers to correspond basically to the parameters discussed herein with respect to liposome preparations. Selecting and providing such parameter based inter alia on straightforward experimentation, is well within the skill of an ordinary worker experienced in this art.

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- A presently highly preferred use of the inventive liposome preparations is in the treatment of infections of the lower respiratory tract, including trachea, bronchi and alveoli, especially when the liposome preparations contain povidone iodine. Also in this indication, the inventive antiseptic preparations, especially those
- 5 containing PVP iodine, have the great advantage of not causing resistances and lead to much less allergic reactions, while permitting a very cost-efficient therapy with a broad spectrum of effect. A povidone iodine liposome preparation according to this invention is e.g. effective against viruses. Further, a liposome preparation of a microbicidal agent such as povidone iodine provides protracted
- 10 release of the agent from liposomes delivering the agent to the pulmonary regions, for example to the alveolar regions of the lung. This leads to extended effect of the antimicrobial substance, and thus less frequent application, as compared with the customary antiseptic solution preparations.
- 15 The present invention is also useful in the treatment of infectious diseases or for alleviation of diseases such as HIV infections which are accompanied by opportunistic infections. Also patients having a suppressed immune system, for example, after organ transplants, can be treated according to the invention. In particular, acute and chronical bronchitis, pneumonia, bronchiectasia, cystic
- 20 fibrosis, diphtheria, tuberculosis can be treated with the povidone iodine preparation according to the invention.

Further highly preferred use is in tissue repair, especially in functional and cosmetic tissue remodelling.

25

Preparations according to this invention can take a variety of forms, which are suitable for administration via the lower respiratory tract, including pharmaceutically acceptable solid or liquid formulations, which are suitable for the generation of inhalable particles. Preparations according to this invention can be

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therefore in the form of (powder) aerosol or in the form of a compacted solid medicament reservoir, preferably a ring tablet, more preferably a gelatine capsule, a powder, a spray, an emulsion, a dispersion, a suspension or even a solution containing the carrier and agent or agents.

5

Generally, the amount of active agents in an inventive preparation will be determined by the desired effect, on the one hand, and the carrying capacity of the carrier preparation for the agent, on the other hand.

10 For inventive preparations with large amounts of active agents or high dosages of active agent, nebulized preparations or aerosols are preferred to powders or powder aerosols. Broadly, the amount of active agent in an inventive carrier preparation can range in concentrations between the lower limit of effectiveness of the agent and the maximum loading of the agent in the respective carrier
15 preparation.

More specifically, for an antiseptic agent, such as povidone iodine, a solution or dispersion in an inventive carrier preparation, especially where the carrier is a liposome preparation, can contain between 0.1 and 10 g of agent in 100 g of
20 preparation. Such a preparation will then typically contain between 1 and 5 g of liposome membrane-forming substance, especially lecithin, per 100 g of preparation.

An inventive aerosol or spray preparation will often comprise up to 50 mg, but
25 could comprise up to and above 100 mg of liposomal active agent formulation and can, for example, be administered by 5 spray doses, each containing 20 mg of liposomal active agent formulation.

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The preparation will typically comprise at least 10 % wt of active agent such as PVP-iodine in the loaded liposomes (or alternative carrier particles), but may comprise up to 50 wt.-% or even more of active agent. Where the active agent is PVP-iodine, the amount of available iodine will generally be about 10 wt.-%
5 (based on PVP-iodine).

More specific formulations are notable from the embodiment examples.

The features and advantages of this invention will become notable in more detail
10 from the ensuing description of preferred embodiments. In these embodiments, which include a best mode, povidone iodine is exemplified as an antiseptic agent and liposomes are chosen as the carrier. This should, however, not be construed as a restriction of this invention to antiseptic agents or, among antiseptic agents, to povidone iodine, and/or to liposomes as the carrier, although such preparations are
15 specifically preferred.

One preferred method for producing the invention's liposomes can generally be described as follows:

20 The lipid membrane-forming components, e.g. lecithin, are dissolved in a suitable solvent such as chloroform or a 2:1 mixture of methanol and chloroform and are filtered under sterile conditions. Then, a lipid film is produced on a sterile high surface substrate, such as glass beads, by controlled evaporation of the solvent. In some cases, it can be quite sufficient to form the film on the inner surface of the
25 vessel used in evaporating the solvent, without using a specific substrate to increase the surface.

An aqueous system is prepared from electrolyte components and the (one or more) active agents to be incorporated in the liposome preparation. Such an aqueous

system can e.g. comprise 10 mmol/l sodium hydrogen phosphate and 0.9 % sodium chloride, at pH 7.4; the aqueous system will further comprise at least the desired amount of the active agent, which in the embodiment examples is povidone iodine. Often, the aqueous system will comprise an excess amount of agent or agents.

The liposomes are generally formed by agitating said aqueous system in the presence of said film formed by the lipid components. At this stage, further additives can be added to improve liposome formation; e.g. sodium cholate can be added. Liposome formation can also be influenced by mechanical action such as pressure filtration through e.g. polycarbonate membranes, or centrifuging. Generally, the raw liposome dispersion will be washed, e.g. with electrolyte solution as used in preparing the above-described solution of the active agent.

When liposomes with the required size distribution have been obtained and washed, they can be redispersed in an electrolyte solution as already described, often also comprising sugars such as saccharose or a suitable sugar substitute. The dispersion can be freeze-dried, and it can be lyophilysed. It can, prior to use, be reconstituted by addition of water and suitable mechanical agitation at the transition temperature of the lipid component, which for hydrogenated soy bean lecithin is e.g. 55°C.

In the following Examples, hydrogenated soy bean lecithin (EPIKURON (TM) 200 SH obtainable from Lukas Meyer, Germany or PHOSPOLIPON (TM) 90 H obtainable from Nattermann Phospholipid GmbH, Germany) was used. However, other pharmaceutically acceptable liposome membrane-forming substances can be used instead, and the person skilled in the art will find it easy to select suitable alternative liposome forming systems from what is described in prior art.

Embodiment Example I

In a 1000 ml glass flask, provided with glass beads for increased surface, 51.9 mg cholesterol and 213 mg hydrogenated soy bean lecithin were dissolved in a
5 sufficient amount of a mixture of methanol and chloroform in a 2:1 ratio. The solvent was then evaporated under vacuum until a film was formed on the inner surface of the flask and on the glass beads.

2.4 g PVP iodine (containing about 10 % available iodine) were separately
10 dissolved in 12 ml water.

Again in a separate vessel, 8.77 g sodium chloride and 1.78 g $\text{Na}_2\text{HPO}_4 \cdot 2\text{H}_2\text{O}$ were dissolved in 400 ml water. Further water was added up to a total volume of 980 ml, and then, approximately 12 ml 1N hydrochloric acid were added to adjust
15 pH to 7.4. This solution was then topped up with water to exactly 1000 ml.

In a fourth vessel, 900 mg saccharose and 57 mg disodium succinate were dissolved in 12 ml water.

20 The PVP iodine solution was then added to the lipid film in the flask and the mixture was shaken until the film dissolved. The resulting liposome formulation was separated from the hydrated lipids in the flask. The product was centrifuged and the supernatant liquid was discarded. The saccharose solution was added ad 12 ml and the product was again centrifuged. Afterwards the supernatant liquid
25 was again discarded. At this stage, a further washing step, using the saccharose solution or the sodium chloride buffer solution could be carried out.

After the last centrifugation step and discarding of the supernatant, 12 ml sodium chloride buffer solution was added, and the liposomes were homogenously

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distributed therein. The product was then distributed into vials each containing 2 ml liposome dispersion, and the vials were then subjected to a freeze-drying step.

After the freeze-drying, each vial comprised about 40 mg solids.

5

The method of Embodiment Example I has a minor disadvantage in that the PVP iodine solution used, due to the high percentage of solids, is rather viscous and thus more difficult to handle.

10 Embodiment Example II

In a 2000 ml flask provided with glass beads to increase surface, 173 mg hydrogenated soy bean lecithin and 90 mg disodium succinate were dissolved in approximately 60 ml of a methanol/chloroform mix in a 2:1 ratio. The solvent
15 was removed under vacuum until a film was formed.

4 g PVP iodine (10 % available iodine) were dissolved in 40 ml of the sodium chloride buffer solution described in Embodiment Example I, and were added to the lipid film in the flask. The flask was then shaken until the film dissolved and
20 liposomes were formed.

The product was centrifuged and the supernatant liquid was discarded.

To the thus produced liposome pellet, further 40 ml sodium chloride buffer
25 solution was added, and the centrifuging step was repeated. The supernatant was again discarded. At this stage, the washing step could be repeated where necessary.

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After the final centrifuging and decanting step, 40 ml sodium chloride buffer solution was again added to the precipitated liposomes. The homogenous dispersion was then distributed into vials, each vial containing about 2 ml liposome dispersion, and the vials were then subjected to a freeze-drying step.

5 This produced approximately 200 mg freeze-dried solids per vial.

Like that of Embodiment Example I, the above-described method uses a hydrating step after film formation in the presence of organic solvents and aims at inclusion rates of 5 to 15 %. These methods generally produce rather large and often multi-

10 lamellar liposomes.

The above-described methods can be modified by a high pressure filtering step through a suitable membrane such as a polycarbonate membrane after the raw liposomes have been formed or after any of the subsequent washing steps or

15 directly by using high pressure homogenisation. This produces much smaller, unilamellar liposomes at increased amounts of encapsulated agent.

Instead of high pressure homogenisation, other prior art methods known to provide small uniform sized liposomes can be employed.

20

Embodiment Example III

A gelatine capsule, which is suitable for the generation of inhalable particles, was prepared from 20 g of povidone iodine liposomes containing leophilised (?????)

25 material according to the above-mentioned general preparation method and 20 mg lactose by applying pressures of up to 500 MPa. From the obtained hard capsule a powder or powder aerosol was generated by abrading methods using a powder inhaler (Orbital-Inhaler by Brin Tech International Ltd.

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It is also possible to prepare embodiments similar to those described above, which comprise an agent capable of promoting the healing of wounds instead of, and not in addition to, the antiseptic agent, such as e.g. povidone iodine disclosed in the above embodiment examples. Presently, it is however preferred to use a
5 wound healing promoting agent (if at all) in addition to an antiseptic agent.

For application of the inventive preparations to a patient, known systems can be used, such as inhalers, powder inhalers, two-chamber gas pressure packs, aerosol spray dispensers, nebulizers, compressors, etc.

10

Embodiment Example IV

Liposomic preparations were aerosolized via an air-driven nebulizer. The output and aerosol characteristics of liposomes with the nebulizer have been previously
15 described. The resulting droplets had a mass medium aerodynamic diameter of about 2.4 μm and are therefore suitable for deposition in the alveolar region.

Using inventive preparations efficiency tests were then carried out, as follows:

20

Test I

This was an in-vitro-test of the bactericidal effect provided by an inventive povidone iodine liposome preparation. The test was based on the quantitative suspension test as described in "Richtlinien der Deutschen Gesellschaft für Hygiene und Mikrobiologie", 1989. In this test, the bactericidal agent is used to
25 kill staphylococcus aureus (ATCC 29213), a major problem in hospital hygiene.

The liposome preparation used was that of Embodiment Example I. At different contact times between 1 und 120 minutes, the minimum concentration of the preparation in water was determined which was capable of killing the

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staphilococci.

The results are shown in Table 1.

5

TABLE I

	<u>Contact Time (Minutes)</u>	<u>Bactericidal Concentration</u>
	1, 2, 3, 4	$\geq 0.060 \%$
	5, 30, 60	$\geq 0.015 \%$
10	120	$\geq 0.007 \%$

The results show that at short contact times (between 1 and 4 minutes) the bactericidal concentration is as low as 0.06 % and that at long contact times (120 minutes) the bactericidal concentration can be as low as 0.007 %.

15

Test II

The virucidal and chlamydicidal activity of liposomal PVP-iodine has been studied, in cell cultures, by Wutzler et al., 9th European Congress for Clinic
20 Microbiology and Infection Diseases, Berlin, March 1999. In cell cultures, liposomal PVP-iodine is highly effective against herpes simplex virus type 1 and adenovirus type 8, while the long-term cytotoxicity experiments indicated that the liposomal form is better tolerated than aqueous PVP-iodine by the majority of cell lines tested. PVP-iodine in liposomal form is not genotoxic.

25

Test III

A 3% PVP-iodine hydrogel liposomal preparation was compared with a 3% PVP-iodine ointment, where the active agent was not in liposomal form. The agent

- 20 -

was applied to standardized in vitro cultures of rat skin and peritoneal explants, as a screening for tissue compatibility of skin and wound antiinfectives.

5 The growth rate of the cultured explants was studied after 30 minutes exposure and incubation with a test substance.

Again, the substantially better toleration of the liposomal preparation was clearly shown in the results, in terms of peritoneum growth rate and skin growth rate.

10 With the ointment, the peritoneum growth rate reached 85%, and the skin growth rate reached 90%; with the liposomal hydrogel formulation, the peritoneum growth rate was 96%, and the skin growth rate was 108%; these values are to be compared with 100% values in a control test using Ringer's solution as the agent.

15

Claims

1. A process for the manufacture of a pharmaceutical preparation for the application of antiseptic agents and/or agents which promote the healing of wounds to the lower respiratory tract, characterised in that the preparation contains at least one of said agents combined with a particulate carrier.
2. The process of claim 1, characterised in that said particulate carrier comprises at least one of a liposome preparation, a microsphere preparation, a nanoparticle preparation, a Large Porous Particle preparation or a laser-pulse polymer coated molecule preparation.
3. The process according to claim 1 or 2, characterised in that at least the greatest part of said agent is encapsulated inside the carrier, especially a liposome or microsphere carrier.
4. The process of any one of claims 1 to 3, characterised in that the antiseptic agent is selected from oxygen- and halogen-releasing compounds; metal compounds, such as silver and mercury compounds; organic disinfectants including inter alia formaldehyde-releasing compounds, alcohols, phenols including alkyl- and arylphenols as well as halogenated phenols, quinolines and acridines,

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hexahydropyrimidines, quaternary ammonium compounds and iminium salts, and guanidines.

5. The process according to claim 4,
5 characterised in that the antiseptic agent is selected from the group comprising metal compounds such as mercury compounds, phenol derivatives such as thymol, eugenol and hexachlorophene, iodine and iodine complexes.

6. The process according to claim 5,
10 characterised in that the antiseptic agent is povidone iodine.

7. The process according to any one of claims 1 to 6,
characterised in that the wound-healing promoting agent is selected from agents promoting granulation and epithelization such as dexpanthenol, allantoin, 15 azulenes, tannines, compounds from the vitamin B series, or similarly acting agents.

8. The process according to any one of the preceding claims,
characterised in that the preparation contains at least one antiseptic and at least
20 one wound-healing promoting agent.

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9. The process according to any one of the preceding claims, characterised in that the carrier particles, especially liposomes, have a substantially uniform size in the range between about 1 and about 50 μm , preferably in the range between about 1 and about 30 μm .

5

10. The process according to claim 9, characterised in that the carrier particles, especially liposomes, have a substantially uniform size in the range between about 20 and 30 μm diameter for application to the trachea, in the range between about 10 and 20 μm diameter for application to the bronchi and between about 1 and 6 μm , especially between 2 and 5 μm , diameter for application to the alveoli.

11. The process according to any one of the preceding claims, characterised in that the carrier, especially liposome, preparation releases the agent over an extended time period, preferably an extended time period of several hours duration.

12. The process according to claim 11, characterised in that the carrier, especially liposome, preparation releases the agent at approximately the same release rate over the release time period.

20

- 24 -

13. The process according to any one of the preceding claims, characterised in that the preparation additionally comprises at least one anaesthetically active agent.

5 14. The process according to any one of the preceding claims, characterised in that the preparation contains additives and adjuvants such as conserving agents, antioxidants and consistency-forming additives.

10 15. The process according to any one of claims 1 to 14, the preparation being in a suitable form for administration via the lower respiratory tract comprising the active-agent loaded carrier, especially in the form of liposomes, preferably in the form of an aerosol, especially in the form of a powder aerosol.

16. The process according to any one of claims 1 to 14, the preparation
15 being in the form of a compacted solid medicament reservoir, preferably a ring-tablet, more preferably a gelatin capsule, a powder, a spray, an emulsion, a dispersion, a suspension or a solution containing the carrier and agent or agents in a pharmaceutically acceptable solid or liquid formulation, which is suitable for the generation of inhalable particles.

- 25 -

17. The process according to any one of the preceding claims, the preparation being in a suitable form for administration via the lower respiratory tract, which comprises:

- a) liposomes comprising a pharmaceutically acceptable liposome membrane forming substance; and
- b) a 0.1 to 2 % PVP iodine solution (at approximately 10 % available iodine in the PVP iodine complex) at least most of which is encapsulated by said liposome membranes,

wherein the liposomes are of substantially uniform size between about 1 and about 50 μm , and, in case, the formulation additionally comprises customary additives, adjuvants and auxiliary substances of a pharmaceutical formulation.

18. The process according to claim 17, characterised in that the liposomes are of substantially uniform size, in the range between about 20 and 30 μm diameter for application to the trachea, in the range between about 10 and 20 μm diameter for application to the bronchi and between about 1 and 6 μm diameter, preferably between about 2 and 5 μm diameter, for application to the alveoli.

19. The process according to any one of claims 1 to 18, wherein the preparation is suited for the treatment of infectious diseases or alleviation of diseases such as HIV infections which are accompanied by opportunistic infections

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or a suppressed immune system.

20. The process according to any one of claims 1 to 18, wherein the preparation is suited for the treatment of acute and chronic bronchitis, pneumonia,
5 bronchiectasia, cystic fibrosis, diphtheria and/or tuberculosis.

21. The process according to any one of claims 1 to 20, wherein the preparation is suited for functional and cosmetic tissue remodelling and repair treatments.

10

22. A method of preventing or treating infections of the human or animal lower respiratory tract, by applying, to said tract, a pharmaceutical preparation comprising at least one antiseptic agent and/or wound-healing promoting agent, said agent being combined with a particulate carrier in said
15 preparation.

23. A method of functional and cosmetic tissue remodelling and repair in the human or animal lower respiratory tract, by applying, to said tract, a pharmaceutical preparation comprising at least one anti-inflammatory especially
20 antiseptic and/or wound-healing promoting agent combined with a particular carrier.

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24. The method of claim 22 or 23, wherein said carrier comprises at least one of a liposome preparation, a microsphere preparation, a nanoparticle preparation, a Large Porous Particle preparation, or a laser-pulse polymer coated molecule preparation.

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25. The method of claim 22 or 23, wherein at least the greatest part of said agent is encapsulated inside the carrier, especially a liposome or microsphere carrier.

10

26. The method of claim 23, wherein the anti-inflammatory agent is selected from antiseptic agents, antibiotics, corticosteroids and wound-healing promoting agents.

15

27. The method of claim 22 or 23, wherein the antiseptic agent is selected from oxygen- and halogen-releasing compounds; metal compounds, such as silver and mercury compounds; organic disinfectants including inter alia formaldehyde-releasing compounds, alcohols, phenols including alkyl- and arylphenols as well as halogenated phenols, quinolines and acridines, hexahydropyrimidines, quaternary ammonium compounds and iminium salts, and

20 guanidines.

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28. The method of claim 22 or 23, wherein the antiseptic agent is selected from the group comprising metal compounds such as mercury compounds phenol derivatives such as thymol, eugenol and hexachlorophene, iodine and iodine complexes.

5

29. The method of claim 22 or 23, wherein the antiseptic agent is povidone iodine.

30. The method of claim 22 or 23, wherein the wound-healing promoting agent is selected from agents promoting granulation and epithelization such as dexpanthenol, allantoines, azulenes, tannines, compounds from the vitamin B series or similarly acting agents.

31. The method of claim 22 or 23, wherein the preparation contains at least one antiseptic and at least one wound-healing promoting agent.

32. The method of claim 22 or 23, wherein the carrier particles, especially liposomes, have a substantially uniform size in the range between about 1 and about 50 μm , preferably in the range between about 1 and about 30 μm .

20

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33. The method of claim 32, wherein the carrier particles, especially liposomes, have substantially uniform size in the range between about 20 and 30 μm diameter for application to the trachea, in the range between about 10 and 20 μm diameter for application to the bronchi and between about 1 and 6 μm diameter, especially between 2 and 5 μm , for application to the alveoli.

34. The method of claim 22 or 23, wherein the carrier, especially liposome, preparation releases the agent over an extended time period, preferably an extended time period of several hours duration.

10

35. The method of claim 22 or 23, wherein the carrier, especially liposome, preparation releases the agent at approximately the same release rate over the release time period.

36. The method of claim 22 or 23, wherein the preparation additionally comprises at least one anaesthetically active agent.

37. The method of claim 22 or 23, wherein the preparation contains additives and adjuvants such as conserving agents, antioxidants and consistency-forming additives.

20

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38. The method of claim 22 or 23, the preparation being in a suitable form for administration via the lower respiratory tract comprising the active-agent loaded carrier, especially in the form of liposomes, preferably in the form of an aerosol, especially in the form of a powder aerosol.

5

39. The method of claim 22 or 23, the preparation being in the form of a compacted solid medicament reservoir, preferably a ring-tablet, more preferably a gelatine capsule, a powder, a spray, an emulsion, a dispersion, a suspension or a solution containing the carrier and agent or agents in a pharmaceutically acceptable solid or liquid formulation, which is suitable for the generation of inhalable particles.

10

40. The method of claim 22 or 23, the preparation being in a suitable form for administration via the lower respiratory tract, which comprises:

15

a) liposomes comprising a pharmaceutically acceptable liposome membrane forming substance; and

b) a 0.1 to 2 % PVP iodine solution (at approximately 10 % available iodine in the PVP iodine complex) at least most of which is encapsulated by said liposome membranes,

20

wherein the liposomes are of substantially uniform size between about 1 and about 50 μm , and, in case, the formulation additionally comprises customary additives, adjuvants and auxiliary substances of a pharmaceutical formulation.

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41. The method of claim 22 or 23, wherein the liposomes are of substantially uniform size, between about 20 and 30 μm diameter for application to the trachea, between about 10 and 20 μm diameter for application to the bronchi and between about 1 and 6 μm , preferably between about 2 and 5 μm diameter, for application to the alveoli.

42. The method of claim 22 or 23, wherein the preparation is suited for the treatment of infectious diseases or alleviation of diseases such as HIV infections which are accompanied by opportunistic infections or a suppressed immune system.

43. The method of claim 22 or 23, wherein the preparation is suited for the treatment of acute and chronic bronchitis, pneumonia, bronchiectasia, cystic fibrosis, diphtheria and/or tuberculosis.

DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that

My residence, post office address and citizenship are as stated below next to my name

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

PREPARATIONS FOR THE APPLICATION OF ANTI-INFLAMMATORY, ESPECIALLY ANTISEPTIC AGENTS AND/OR AGENTS PROMOTING THE HEALING OF WOUNDS, TO THE LOWER RESPIRATORY TRACT the specification of which (check one)

☒ is attached hereto

☐ was filed on 27 May 1999 as Application Serial No. PCT/EP99/03681 and was amended on _____ (if applicable)

I hereby authorize and request our attorney, Davidson, Davidson & Kappel, LLC of 1140 Avenue of the Americas, New York, New York 10036 to insert here in parentheses (Application number _____, filed _____) the filing date and application number of said application when known

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose all information which is known to me to be material to the patentability of this application as defined in Title 37, Code of Federal Regulations, §1.56

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign and/or provisional application(s) for patent or inventor's certificate listed below and have also identified below any foreign and/or provisional application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed

PRIOR APPLICATION(S)

<u>60/086,895</u>	<u>U.S.</u>	<u>27 May 1998</u>	<input checked="" type="checkbox"/> <u>27 May 1998</u>	<input checked="" type="checkbox"/>
(Number)	(Country)	(Day/Month/Year Filed)	Yes	No
_____	_____	_____	Yes	No
(Number)	(Country)	(Day/Month/Year Filed)	Yes	No

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, §1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application

_____	_____	_____
(Application Serial Number)	(Filing Date)	(Status) (patented, pending, abandoned)
_____	_____	_____
(Application Serial Number)	(Filing Date)	(Status) (patented, pending, abandoned)

8 And I hereby appoint Clifford M. Davidson, Registration No. 32,728, Leslye B. Davidson, Registration No. 38,854, Cary S. Kappel, Registration No. 36,561, Morey B. Wildes, Registration No. 36,968, Robert J. Paradiso, Registration No. 41,240, Scott L. Appelbaum, Registration No. 41,587, Cynthia R. Moore, Registration No. 46,086 and David G. Knasiak, Registration No. 45,991 my attorneys, with full power of substitution and revocation, to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith, correspondence address DAVIDSON, DAVIDSON & KAPPEL, LLC, 1140 Avenue of the Americas, 15th Floor, New York, New York 10036, Telephone (212) 997-1028; Fax: (212) 997-1037

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon

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